CLAIMS

- 1. A vaccine composition comprising at least one group A Streptococcal antigen and a proteosome adjuvant.
 - 2. A vaccine composition according to claim 1 wherein the antigen comprises an S. pyogenes M protein peptide.
- 3. A vaccine composition according to claim 1 wherein the antigen comprises a fragment of the M protein from the C-terminal region of between 6 and 25 amino acids in length.
- 4. A vaccine composition according to claim 1 wherein the antigen has
 the sequence ASREAKKQVEQKALE.
 - 5. A vaccine composition according to claim 2 wherein the antigen is flanked by amino acid sequences to maintain helical folding of the antigen.
- 20 6. A vaccine composition according to claim 5 wherein the peptide antigen has the sequence KQAEDKVKASREAKKQVEKALEQLEDKVK.

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- 7. A vaccine composition according to claim 1 wherein the antigen is selected from an MtsA peptide or a protein H peptide.
- 8. A vaccine composition according to claim 1 wherein the antigenic peptide further comprises a C- or N-terminal hydrophobic moiety for insertion of the antigenic pepetide into proteosome adjuvant vesicles.
- 9. A vaccine composition according to claim 1 for parenteral or oral administration.

- 10. A vaccine composition according to claim 9 for intranasal administration.
- 5 11. A vaccine composition according to claim 1 for use in the treatment or prophylaxis of a group A Streptococcal infection in an individual.
 - 12. A vaccine composition according to claim 11 wherein the composition is administered intranasally to the individual.

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- 13. A vaccine composition according to claim 11 wherein administration of the composition to the individual induces a mucosal immune response.
- 14. A vaccine composition according to claim 11, wherein administration of the vaccine composition induces a serum immune response.
 - 15. A vaccine composition according to claim 11 wherein the vaccine composition is administered intranasally to the individual and treatment or prophylaxis of group A Streptococcal infection is produced via reduction or prevention of bacterial colonisation of the throat.
 - 16. A method of treatment of prophylaxis of group A Streptococcal infection in an individual comprising administering a vaccine composition according to claim 1 to the individual.

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- 17. A method according to claim 16 wherein said vaccine composition is administered intranasally to said individual.
- 18. A method according to claim 17 wherein the treatment or prophylaxis
 group A Streptococcal infection is produced via prevention or reduction of bacterial
 colonisation of the throat.